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(54) Pharmaceutical compositions containing insulin.

(5) The invention provides a pharmaceutical composition for the oral administration of insulin comprising insulin, a bile acid or alkali metal salt thereof, the bile acid being selected from the group consisting of cholic acid, chenodeoxycholic acid, taurocholic acid, taurochenodeoxycholic acid, glycocholic acid, glycochenocholic acid, 3 $\beta$ -hydroxy-12-ketocholeic acid, 12 $\alpha$ -3 $\beta$ -dihydrocholeic acid, and ursodesoxycholic acid, and a protease inhibitor, the composition being provided with an enterocoating to assure passage through the stomach and release in the intestine.

PHARMACEUTICAL COMPOSITIONS CONTAINING INSULIN

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The present invention relates to a pharmaceutical composition containing insulin. More particularly, the present invention relates to a pharmaceutical composition 5 for the oral administration of insulin.

Insulin is a medicament particularly useful as a hypoglycaemic agent being widely used by patients suffering from diabetes and is the only treatment for juvenile diabetes 10 mellitus.

In practice today insulin is administered only by injection. The everyday injection of insulin is very troublesome and causes considerable physical and even mental suffering 15 for the patients. Several severe side effects such as lipodystrophy at the site of the injection, lipoatrophy, lipohypertrophy or occasional hypoglycemia have been noted and reported to occur.

To avoid the daily injection of the drug, the insulin pump 20 has been developed in the last decade. This pump, however, also suffers from some of the disadvantages of the daily injection. Since insulin is normally secreted into the portal vein, normally the liver is exposed to a greater insulin concentration than peripheral tissues. Insulin administered via 25 the peripheral venous system to insulin-deficient diabetic patients results in a concentration of insulin in the portal vein almost equal to that in the peripheral circulation. The net result is hypoinsulinemia in the portal vein and the liver and 30 hyperinsulinemia in the peripheral venous system. This is due to

In order to overcome the difficulties caused by ~~injection~~ **Q127535** of insulin, rectal administration of insulin has recently been proposed, studied and developed.

5 Shichiri et al (J. Pharm. Pharmac. 30, 806-808, 1978),  
Bar-On et al (Br. J. Pharmac. 73, 21-24, 1981), and others  
10 tested the hypoglycaemic effect of insulin mixed with poly-  
oxyethylen lauryl ether or polyoxyethylene-20-cetyl ether by  
administering through the rectum. Ziv et al (Life Sciences, 29,  
15 803-809, 1981) tested the same effect with insulin mixed with  
bile salts. The insulin affected the blood glucose levels, by  
reduction of approximately 50%, with dose of 48 $\mu$ /kg.

20 In a further article by Ziv, Kidron, Bar-On and Berry  
(Life Sciences, 31, pp. 2837-2841, 1982) insulin was used as  
a model for proteins in general to discover the theoretical  
question of protein absorption through the intestine and it  
was found that in the presence of the strong detergent effect of  
25 deoxycholic acid and soybean trypsin inhibitor, biologically  
active macromolecules such as insulin could be effectively  
absorbed from the intestine.

30 Similarly, in British Patent 1,563,311 there is described  
and claimed a pharmaceutical composition for rectal administra-  
tion which comprises insulin, a carrier suiting the composition  
for rectal administration, and an agent for increasing the rate  
of absorption of the insulin into the body on rectal adminstration  
of the composition, the agent comprising at least one material

number of polyoxyethylene units is 4 to 30, (b) anionic surface  
surface active agents, (c) bile acids and (d) alkali metal salts

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of bile acids and amounting to 0.001 to 0.5 times the weight of the carrier. In U.S. Patents 4434159 and 4164573 there are described similar insulin containing pharmaceutical compositions for rectal administration.

Thus the administration of insulin through the portal system of the human rectum in suppository form or further along the intestinal tract, e.g., by enema-like introduction is suggested and taught by said articles and patent.

Nevertheless it has been found that only part of the insulin is absorbed through the portal system from the human rectum and rectal administration also represents a major inconvenience for the patient.

According to the present invention, there have now been developed pharmaceutical compositions for administering insulin which overcome all of the above-mentioned disadvantages of the prior art systems.

More specifically, there have now been discovered and provided according to the present invention pharmaceutical compositions for the oral administration of insulin comprising insulin, a bile acid or alkali metal salt thereof, said bile acid being selected from the group consisting of cholic acid, chenodeoxycholic acid, taurocholic acid, taurochenodeoxycholic acid, glycocholic acid, glycochenocholic acid,  $3\beta$ -monohydroxychloric acid, lithocholic acid,  $3\alpha$ -hydroxy-12-ketocholic acid,  $3\beta$ -hydroxy-12-ketocholic acid,  $12\alpha$ - $3\beta$ -dihydrocholic acid, and ursodesoxycholic acid, and a protein, insulin.

Thus there have now been discovered pharmaceutical compositions containing insulin which can be administered orally and which have the same effect as naturally secreted insulin on the blood glucose levels. The insulin administered according to the present invention reaches the intestine and is quickly absorbed in the body through the intestine and through the portal system to the liver. This absorption route is the most convenient way to administer the drug and it resembles the physiological secretion of insulin by the pancreas, thus enabling delicate control of the blood glucose level and the metabolic activities of the liver and the peripheral organs controlled by insulin.

Various attempts have been made in the past to administer insulin orally. In one study it was shown that administration of liposome-entrapped insulin caused a significant reduction of blood glucose levels in diabetic rats (Dapergolas, G. and Gregoriadis, Lancet ii, 824-827, 1976). Patel and Ryman (FEBS Letters, 62, 60-63, 1976) showed that insulin administered orally entrapped in liposomes is effective in diabetic rats. Papahadjopoulos and Sjoka (U.S. patent No. 4,235,871) suggested to use liposomes to encapsulate insulin and Sears (U.S. patent No. 4,145,410) used synthetic phosphatidyl compounds to stabilize the liposomes against lipolysis.

Another approach for insulin enhanced activity is the addition of an adjuvant such as choline (which is not a bile salt) to the insulin injections (U.S. patent 2563070). This is totally different

from the present invention in that the adjuvant is administered from the intestinal lumen to the blood circulation while with

injectable solutions no such absorption takes place or is necessary and the function of choline which is different structurally and chemically from cholic acid is entirely 5 different in said patent and is intended to delay the insulin absorption.

Thus it will be realized that none of the said publications 10 teaches or suggests the novel pharmaceutical composition of the present invention which includes the use of bile salts to promote the absorption of insulin, the use of protease inhibitors to protect insulin against proteolysis and the use of enterocoating of the active mixture.

15 Human insulin including human insulin genetically reproduced or any insulin such as, for example, the insulin obtained from cows (bovine), pigs or whales can be used as the insulin for compositions of this invention. Furthermore, 20 metal complexes of insulin such as the zinc complex of insulin as well as protamine zinc insulin and globin zinc insulin may be also used as the insulin in compositions of this invention.

25 The protease inhibitor used in the compositions of the present invention can be any material which has the ability to inhibit any proteolytic activity.

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Practical examples of such protease inhibitors include aprotinin (Trasylol (R) of Bayer), Pentamidine isethionate, 5 antipain, tosylamide-phenylethyl-chloromethyl ketone (TPCK), phenylmethyl sulfonylfluoride (PMSF), pepstatin, trypsin inhibitor, Acetone, Alcohols, guanidium,  $A_2$ -macroglobulin, TLCK, Chelating agents of Zn, Iodoacetate,  $\alpha_1$ -antitrypsin, EDTA, Zn, Antithrombin III, leupeptin, Trypsin inhibitor from soy bean, trypsin inhibitor from 10 hen egg white, trypsin inhibitor from chicken egg white, etc.

Some of the above protease inhibitors might be toxic in large doses and therefore, if chosen, the use and dosage thereof must be carefully screened and tested.

15 In especially preferred embodiments of the present invention said protease inhibitor is selected from the group consisting of aprotinin,  $A_2$ -macroglobulin, antithrombin III and trypsin inhibitor from soy bean or chicken egg white.

20 The most preferred protease inhibitor agents used in this invention are preferably Trasylol (R) in the amount of 1000 k.i.u./ 100 mg pill, or 3 mg soybean trypsin inhibitor or 10 mg soybean flour.

25 The above-mentioned bile acids and alkali metal salts thereof used in the oral compositions of the present invention promote the absorption of the insulin from the intestinal tract and act as carriers therefor, however, it was interesting and surprising to note that deoxycholic acid, which was the acid of choice in 30 the article in Life Sciences, Vol. 31, pp. 2837-2441 (1982) is

intestinal wall

It has also been surprisingly found that sodium cholate can simultaneously function both as the bile acid carrier of the insulin and the protease inhibitor agent and thus a composition comprising insulin and sodium cholate in an enteric coating is especially preferred.

The amount of insulin in a composition is 20-50u/kg in rats and expected to be about 0.5-3u/kg in humans. Preferred dosages for humans are about 1-2u/kg/treatment with three treatments a day, however sustained release microencapsulation could allow treatment to be reduced to once or twice a day.

The enterocoating and possible microencapsulation of the mixture provides protection for the insulin against decomposition in the stomach and for the slow release of the mixture constituents in the intestinal tract.

The enterocoating is carried out by methods known per se in the art, e.g., according to Remington Pharmaceutical Sciences, p. 1614-1615 (1975, 15th Ed. Mack Pub. Co.) and Theory and Practice of Industrial Pharmacy, Lachman, Liberman & Canig, p. 116-117, 371-374 (1976, 2nd Ed.) as is the enteric micro-encapsulation (Theory and Practice of Industrial Pharmacy ibid, pp. 420-438).

One of the findings of the present invention is that there is different rate of absorption of the different constituents of the present composition from the intestinal lumen into the blood stream. The absorption of the bile acid is very fast.

60 minutes.

For this reason a drug regimen involving ingestion of a pair of pills at spaced intervals, e.g., a second pill containing a higher concentration of bile acid to be taken half an 5 hour after the first pill is contemplated as is microencapsulation of different constituents with spaced time release coatings to enhance the absorption of the insulin into the system.

10 While the invention will now be described in connection with certain preferred embodiments in the following examples it will be understood that it is not intended to limit the invention to these particular embodiments. On the contrary 15 it is intended to cover all alternatives, modifications and equivalents as may be included within the scope of the invention as defined by the appended claims. Thus, the following examples which include preferred embodiments will serve to illustrate the 20 practice of this invention, it being understood that the particulars shown are by way of example and for purposes of illustrative discussion of preferred embodiments of the present 25 invention only and are presented in the cause of providing what is believed to be the most useful and readily understood description of formulation procedures as well as of the principles and conceptual aspects of the invention.

30 EXAMPLE 1.

An enterocoated capsule was prepared for oral administration

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TABLE I

	Composition of this invention	Test No.	Formulation	Time (Min)	Glucose mm	Insulin μm/ml
5		1.	insulin 7.3u/kg	0	36.7	15
	Oral administration		cholate 30 mg	15	33.9	0
			soybean -	30	34.4	6
10			-trypsin inhibitor	45	34.6	6
			<u>20 mg</u>	60	33.8	12
			in two capsules	75	29.1	23
			(enterocoated)	105	23.7	36
				135	21.8	22
15				165	16.1	3
				210	15.8	0
				240	11.5	0
				270	12.1	0
20				300	10.2	0
				330	7.5	0
				360	9.3	0

COMPARISON TABLE II

	Test No.	Formulation	Time (Min.)	Glucose mm	Insulin $\mu$ u/ml
5	10	Insulin 0.5u/kg intramuscular injection	0	28.7	0
			10'	23.3	73
			20'	21.5	76
			30'	19.9	213
			45'	15.6	220
			60'	12.9	81
			75'	10.9	120
			90'	8.5	61
			120'	6.5	50
			150'	6.4	18
			180'	6.5	25

A solution was prepared for direct intestinal administration of 0.5 ml in final volume.

5 In the rat.

TABLE III

	Test No.	Formulation	Percent decrease in blood glucose			No. of Animals tested.
			60 min	120 min	240 min	
10 15 20 25 30 35	1.	Insulin 24u/kg sodium cholate 1% Trasylol 3000 K.I.U. in saline	-33	-11	+11	5
	2.	Insulin 24u/kg sodium cholate 1% in saline	-23	-19	+10	4
	3.	Insulin 24 u/kg sodium cholate 0.5% Trasylol 3000 K.I.U. in saline	-18	-24	-2	6
	4.	Insulin 48u/kg sodium cholate 1% in saline	-34	-20	+10	10
	5.	Insulin 48u/kg sodium cholate 1% Trasylol 1000 u in saline	-55	-50	+3	6
	6.	Insulin 48u/kg sodium cholate 1% Trasylol 3000u in saline	-61	-66	-25	6
	7.	Insulin 48u/kg sodium cholate 1% soybean trypsin	-35	-28	+14	6
Insulin 48u/kg sodium taurocholate 1% in saline			-?	-?	+27	?

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TABLE III (Continued)

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5	Test No.	Formulation	Percent decrease in blood glucose			No. of Animals tested
			60 min	120 min	240 min	
10	10.	Insulin 48 u/kg sodium taurocholate 1% Trasylol 3000 K.I.U. in saline	-39	-36	+3	7
15	11.	Insulin 48u/kg Sodium taurocholate 1% Soybean trypsin-inhibitor 3 mg in saline	-15	-12	+21	6
20	12.	Insulin 48u/kg Trasylol 3000 K.I.U. in saline	-33	-30	+4	6
Control-intestinal administration	13.	Insulin 48u/kg in saline	-5	-6	+25	6

As is clear from the tables the effect of intestinal administration of insulin on blood glucose levels is similar to the effect of insulin injected to the animals. The effect is 5 similar when insulin is given orally to the dog or directly into the intestine of the rat.

Enterocoating provides the sufficient shelter against the destruction of the insulin in the stomach and delays its effect 10 for one hour in the dog.

Now the following examples illustrate practically the pharmaceutical compositions of insulin for oral use embodying 15 this invention, wherein the dosage of insulin employed are for the human bodies.

All examples are for one pill or one capsule containing total weight of 100 mg. The active compounds will be given in 20 detail. The complimentary weight is of inert compounds like manitol or avicel 101.

Thus the active ingredients and the vehicle for oral administration of compositions according to the present invention 25 are hereinafter set forth in tabular form:

Example	Amount Insulin	Bile Acid/Salt	Protease Inhibitor	Vehicle
5	2.	100 I.U. 15 mg sodium cholate	-	enterocoated capsule
	3.	100 I.U. 15 mg sodium cholate	aprotinin 1000 K.I.U.	enterocoated capsule
	4.	100 I.U. 15 mg. sodium cholate	aprotinin 3000 K.I.U.	enterocoated capsule
10	5.	100 I.U. 15 mg. sodium cholate	5 mg soybean trypsin inhibitor	enterocoated capsule
	6.	100 I.U. 15 mg. sodium cholate	-	enterocoated pills
15	7.	100 I.U. 15 mg. sodium cholate	aprotinin 1000 K.I.U.	enterocoated pills
	8.	100 I.U. 15 mg. sodium cholate	aprotinin 3000 K.I.U.	enterocoated pills
	9.	100 I.U. 15 mg. sodium cholate	5 mg chicken egg white trypsin inhibitor	enterocoated pills
20	10.	100 I.U. 15 mg sodium tauro-cholate	aprotin 1000 K.I.U.	enterocoated capsule
	11.	100 I.U. 15 mg. sodium tauro-cholate	aprotinin 3000 K.I.U.	enterocoated capsule
25	12.	100 I.U. 15 mg. sodium tauro-cholate	5 mg soybean trypsin inhibitor	enterocoated capsule
	13.	100 I.U. 15 mg. sodium tauro-cholate	aprotinin 1000 K.I.U.	enterocoated pills
30	14.	100 I.U. 15 mg. sodium cheno-deoxycholate	aprotinin 1000 K.I.U.	enterocoated capsule
	15.	100 I.U. 15 mg. sodium cheno-deoxycholate	aprotinin 3000 K.I.U.	enterocoated capsule

Example	Amount Insulin	Bile Acid/Salt	Protease Inhibitor	Vehicle
5	17. 100 I.U.	15 mg. sodium cheno- deoxycholate	aprotinin 1000 K.I.U.	enterocoated pills
	18. 100 I.U.	15 mg. sodium cheno- deoxycholate	aprotinin 3000 K.I.U.	enterocoated pills
10	19. 100 I.U.	15 mg. sodium cheno- deoxycholate	5 mg. soybean trypsin inhibitor	enterocoated pills

Examples 20 and 21

The following interocoated tablets were prepared in the following manner:

Component	Example 20	Example 21
Insulin	2 mg	2 mg
Sod. Cholate	15 mg	15 mg
Trasitol	-	1000 U
Lactose Hydrous USP	144 mg	150 mg
Starch NF	36 mg	30 mg
Magnesium Stearate NF	3 mg	3 mg
Eudragit L-100 (Polymer of Acrylic and Methacrylic Acid Esters)	4 mg	4 mg
Talc NF	4 mg	4 mg
Polyethylene Glycol 6000 NF	0.4 mg	0.4 mg
25	Total	208.4 mg
		208.4 mg

Method of Preparation:

(a) In order to homogeneously disperse the active components triturations with a solid vehicle such as lactose for example, for each component were individually prepared. Gradual dry mixing of all the components was then performed. The components are then mechanically pressed to form tablets of 9 mm diameter;

(b) A solution of the enterocoating polymer is then prepared by solving the polymer in an organic solvent such as for example a methylene chloride + isopropyl alcohol mixture. The tablets are coated by spraying the solution within a mildly warmed jar while the tablets roll

The dissolution of the tablets was then tested according to USP XX. The tablets were found to be stable for two hours in gastric fluid. When they are then trans-

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It will be evident to those skilled in the art that the invention is not limited to the details of the foregoing illustrative embodiments and examples and that the present invention may be embodied in other specific forms without departing from the essential attributes thereof, and it is, therefore, desired that the present embodiments and examples be considered in all respects as illustrative and not restrictive, reference being made to the appended claims, rather than to the foregoing description, and all changes which come with the meaning and range of equivalency of the claims are, therefore, intended to be embraced therein.

1. A pharmaceutical composition for the oral administration  
5 of insulin comprising insulin, a bile acid or alkali metal  
salt thereof, said bile acid being selected from the group  
consisting of cholic acid, chenodeoxycholic acid, taurocholic  
acid, taurochenodeoxycholic acid, glycocholic acid, glyco-  
10 chenocholic acid, 3 $\beta$ -monohydroxychloric acid, lithocholic  
acid, 3 $\alpha$ -hydroxy-12-ketocholic acid, 3 $\beta$ -hydroxy-12-ketocholic  
acid, 12 $\alpha$ -3 $\beta$ -dihydrocholic acid, and ursodesoxycholic acid,  
15 and a protease inhibitor, said composition being provided  
with an enterocoating to assure passage through the stomach  
and release in the intestine.

2. A pharmaceutical composition according to  
20 claim 1, wherein said protease inhibitor is selected from  
the group consisting of aprotinin, A<sub>2</sub>-macroglobulin, antithrombin  
III and trypsin inhibitor from soy bean or chicken egg white.

25 3. A pharmaceutical composition according to  
claim 1, comprising sodium cholate as both the bile acid and  
the protease inhibitor.

30 4. A pharmaceutical composition  
according to claim 1, wherein the components of said  
composition are microencapsulated and enterocoated to provide  
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CLAIMS (for AT)

1. A process for preparing a pharmaceutical composition for the oral administration of insulin, characterized in that it comprises mixing active ingredients consisting of insulin, a bile acid or alkali metal salt thereof, said bile acid being selected from the group consisting of cholic acid, chenodeoxycholic acid, taurocholic acid, taurochenodeoxycholic acid, glycocholic acid, glycochenocholic acid,  $3\beta$ -monohydroxychloric acid, lithocholic acid,  $3\alpha$ -hydroxy-12-ketocholic acid,  $3\beta$ -hydroxy-12-ketocholic acid,  $12\alpha$ - $3\beta$ -dihydrocholic acid, and ursodesoxycholic acid, and a protease inhibitor,  
and providing said active components with an enterocoating to assure passage through the stomach and release in the intestine.
- 10 2. Process according to claim 1, characterized in that said active components are mixed with a solid vehicle, the mixture thus obtained is mechanically pressed to form tablets, the later are coated with a solution of an enterocoating polymer in an organic solvent and the organic solvent is then removed from the coated tablets.
- 15 3. Process according to claim 2, characterized in that the solid vehicle is lactose and the organic solvent is a methylene chloride-isopropyl alcohol mixture.



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(57) The invention provides a pharmaceutical composition for the oral administration of insulin comprising insulin, a bile acid or alkali metal salt thereof, the bile acid being selected from the group consisting of cholic acid, chenodeoxycholic acid, taurocholic acid, taurochenodeoxycholic acid, glycocholic acid, glycochenocholic acid, 3 $\beta$ -hydroxy-12-ketocholic acid, 12 $\alpha$ -3 $\beta$ -dihydrocholic acid, and ursodesoxycholic acid, and a protease inhibitor, the composition being provided with an enterocoating to assure passage through the stomach and release in the intestine.

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DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 2)
A	EP-A-0 071 433 (KOWA CO. LTD.) * Page 36, claim 1 *	1-4	A 61 K 37/26 A 61 K 37/64 A 61 K 31/575 // (A 61 K 37/64 A 61 K 37:26 A 61 K 31:575)
A	FR-M- 662 (HENRY MARINUS CHRISTENSEN) * Page 2, right-hand column - page 3; abstract *	1-4	
A	DE-A-2 640 707 (K. THEURER) * Page 1, claim *	1-4	
A	--- CHEMICAL ABSTRACTS, vol. 93, no. 10, 8th September 1980, page 388, abstract no. 101445z, Columbus, Ohio, US; K. ICHIKAWA et al.: "Rectal absorption of insulin suppositories in rabbits", & J. PHARM. PHARMACOL. 1980, 32(5), 314-18 * Abstract *	1-4	TECHNICAL FIELDS SEARCHED (Int. Cl. 2) A 61 K